To:

KIM, Eui-Bak

The Cheonghwa Bldg., 1571-18, Seocho-dong, Seocho-gu, 137-

874 Seoul, Republic of Korea

DEC 20 2004

PCT

NOTIFICATION OF TRANSMITTAL OF INTERNATIONAL PRELIMINARY **EXAMINATION REPORT** 

(PCT Rule 71.1)

Date of mailing

(day/month/year) 17 DECEMBER 2004 (17.12.2004)

Applicant's or agent's file reference

HL-20133-PCT

IMPORTANT NOTIFICATION

International application No.

International filing date (day/month/year)

ATENT COOPERATION TREATY

Priority date (day/months/year)

PCT/KR2003/001849 08 SEPTEMBER 2003 (08.09.2003) 11 SEPTEMBER 2002 (11.09.2002)

Applicant

HANLIM PHARMACEUTICAL CO., LTD. et al

- 1. The applicant is hereby notified that International Preliminary Examining Authority transmits here with the international preliminary examination report and its annexes, if any, established on the international application.
- 2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
- 3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report(but not of any annexes) and will transmit such translation to those Offices.

#### 4. REMINDER

The applicant must enter the national phase before each elected office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details in the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/KR

Korean Intellectual Property Office 920 Dunsan-dong, Seo-gu. Dacjeon 302-701. Republic of Korea

Facsimile No. 82-42-472-7140

Authorized officer

COMMISSIONER

Telephone No. 82-42-481-5762



### **COPY FOR IB**

Rcc'd PCT/PTO 0 9 MAR 2005

# PATENT COOPERATION TREATY

REC'D 0 3 JAN 2005

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### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

**PCT** 

(PCT Artcle 36 and Rule 70)

Applicant's or agent's file reference HL-20133-PCT	Examination Report (Form PCT/IPEA/416)				
International application No. PCT/KR2003/001849	International filing date(day/mo		Priority date (day/month/year 11 SEPTEMBER 2002 (11.0		
International Patent Classification (IPC)  IPC7 C07D 211/90  Applicant	) or national classification and IF	PC			
HANLIM PHARMACEUTIC	CAL CO., LTD. et al				
and is transmitted to the applica			•	ng Authority	
This report is also accom	and of sheets, including a sheets, including a sheets, i.e., sheets for this report and/or sheets could be administrative Instructions of the Administrative Instructions of the sheets.	ts of the description taining rectification	on, claims and/or drawings wh	nich have been rity (see Rule	
These annexes consist of a total	al ofsheets.				
I X Basis of the report II Priority III Non-establishment IV Lack of unity of the vertex of the vert	nt of opinion with regard to nove invention nent under Article 35(2) with reg lanations supporting such statem	elty, inventive step ard to novelty, inv		ability;	
Date of submission of the demand	I	Date of completion	of this report		
09 MARCH 2004	1 (09.03.2004)	09 DECE	VIBER 2004 (09.12.2004)		
Name and mailing address of the IF  Korean Intellectual Pro  920 Dunsan-dong, Seo Republic of Korea  Facsimile No. 82-42-472-7140	operty Office -gu, Daejeon 302-701,	Authorized officer  KIM, Hee Ji  Telephone No. 8	n '	CIPIN	



### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International aplication No.

PCT/KR2003/001849

I. Basis of the report	
1. With regard to the elements of the international application:*	
X the international application as originally filed	
the description:	, as originally filed
pages	, filed with the demand
pages, filed with the letter of	
the claims:	or anisimally filed
pages, as amended (together win	, as originally filed th any statment) under Article 19
	, filed with the demand
pages, filed with the letter of	
the drawings:	, as originally filed
pages	
pages	
the sequence listing part of the description:	i-ille: Eled
pages	, as originally filed, filed with the demand
pages, filed with the letter of	
With regard to the language, all the elements marked above were available or furnished to this     the international application was filed, unless otherwise indicated under this item.	
These elements were available or furnished to this Authority in the following language	<del></del>
the language of a translation furnished for the purposes of international search (under R	uie 23.1(b)).
the language of publication of the international application (under Rule 48.3(b)).	evenination(under Pules 55 2 and/
the language of the translation furnished for the purposes of international preliminary or 55.3).	examination funder Rules 33.2 and
3. With regard to any nucleotide and/or amino acid sequence disclosed in the international preliminary examination was carried out on the basis of the sequence listing:	application, the international
contained inthe international application in written form.	
filed together with the international application in computer readable form.	
furnished subsequently to this Authority in written form.	
furnished subsequently to this Authority in computer readable form  The statement that the subsequently furnished written sequence listing does not	go beyond the disc losure in the
international applicationas as filed has been furinshed.	
The statement that the information recorded in computer readable form is identical been furnished.	to the written sequence listing has
4. The amendments have resulted in the cancellation of:	
the description, pages	
the claims, Nos.	
the drawings, sheets	
This report has been established as if (some of) the amendments had not been mad go beyond the disclosure as filed, as indicated in the Supplemental Box(Rule 70.2(c))	le, since they have been considered to
* Replacement sheets which have been furnished to the receiving Office in response to an invition in this opinion as "originally filed." and are not annexed to this report since they do not and 70.17).	ation under Article 14 are referred to contain amendments (Rules 70.16
** Any replacement sheet containing such amendments must be referred to under item I and a	nnexed to this report.



#### INTERNATIONAL PRELIMINARY EXAMINATION

International aplication No.

PCT/KR2003/001849

V.	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability;
	citations and explanations supporting such statement

1.	Statement			
	Novelty (N)	Claims	1-11	YES
	• • •	Claims		NO
	Inventive step (IS)	Claims	1-11	YES
		Claims		
	Industrial applicability (IA)	Claims	1-11	
	mountain approximation (in a)	Claims		

2. Citations and explanations (Rule 70.7)

Reference is made to the following document:

D1: WO 95/25722

1. Novelty and Inventive Step

The present invention relates to a process for the preparation of (S)-amlodipine from (R,S)-amlodipine using L-tartaric acid and synthetic intermediates for the preparation of (S)-amlodipine. More specifically, the process of the present invention comprises (i) reacting (R,S)-amlodipine with (L)-tartaric acid in dimethyl sulfoxide, (ii) filtering off the resulting precipitate of step (i), (iii) precipitating (S)-amlodipine-hemi-L-tartarate-DMSO-solvate by adding methylene chloride to the filtrate of step (ii), (iv) optionally forming (S)-amlodipine-hemi-L-tartarate-monohydrate by adding an alcohol to (S)-amlodipine-hemi-L-tartarate-DMSO-solvate obtained in step(iii), (v) treating with a base (S)-amlodipine-hemi-L-tartarate-DMSO-solvate obtained in step(iii) or (S)-amlodipine-hemi-L-tartarate-monohydrate obtained in step (iv).

D1, which is considered to represent the most relevant state of the art, discloses a process for the preparation of (R)- and (S)-isomers of amlodipine from the mixture thereof.

Compared with the present invention, they are the same in that (S)-amlodipine isomer is prepared from the racemic mixture by using tartaric acid as a resolving agent. However, they are different in that the process of D1 employs D-tartaric acid(see example 1,9), while L-tartaric acid which is much cheaper than D-tartaric acid is used in the present invention. Therefore, the process of the present invention is very favorable for industrial-scale mass production. Moreover, it is not obvious to a skilled person in the art to use only L-tartaric acid for the resolution of (S)-amlodipine. Therefore, the present invention is considered to be novel and to involve an inventive step.

2. Industrial Applicability

The present invention is considered to be industrially applicable.



## INTERNATIONAL PRELIMINARY EXAMINATION

International aplication No.

PCT/KR2003/001849

ertain documents cited			
tain published documents (Ru	le 70.10)		
Application No. Patent No.	Publication date (day/month/year)	Filing date (day/month/year)	Priority date (valid claim) (day/month/year)
E,X US 2003/0176706 A1	18/09/2003	18/03/2002	
Document US 2003/0176796 published after the internation	Al was filed on 18/ nal filing date.	03/2002 and published on 18/09/20	003, i.e. filed prior to the priority date bu
on-written disclosures (Rule	70.9)		
ON-Millon disclosures (1701)	, 4.12 )		· .
Kind of non-written di	isclosure I	Date of non-written disclosure (day/month/year)	Date of written disclosure referring to non-written disclosure (day/month/year)
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